What is claimed is:

- 1. A method for the preparation of sevoflurane which comprises:
- (a) providing a liquid mixture of (CF₃)₂CHOCH₂Cl, hydrogen fluoride, and an amine; and
- (b) reacting the mixture; to form (CF₃)₂CHOCH₂F.

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- 10 2. The method of claim 1, wherein the mixture is reacted by heating.
 - 3. The method of claim 1, wherein the mixture is reacted by heating at 40°C to 80°C.
- 4. The method of claim 1, wherein the mixture is reacted by heating at 55°C to 65°C.
- 5. The method of claim 1, wherein the amine is selected from a primary amine, a secondary amine, or a tertiary amine.
 - 6. The method of claim 5, wherein the amine is selected from propylamine or diethylamine.
- The method of claim 5, wherein the amine is a tertiary amine.
 - 8. The method of claim 7, wherein the tertiary amine is a trialkylamine.
- 9. The method of claim 8, wherein the trialkylamine is selected from triethylamine, tripropylamine, triisopropylamine, tributylamine, dimethyl ethyl amine, di-isopropyl ethyl amine, or mixtures thereof.
 - 10. The method of claim 1, wherein the amine is a cyclic amine.
 - 11. The method of claim 10, wherein the cyclic amine is selected from pyrrolidine, N-methyl pyrrolidine, or piperidine.

- 12. The method of claim 1, wherein the mixture comprises (CF₃)₂CHOCHCl₂.
- The method of claim 12, wherein the mixture comprises from 0.01 to 20 percent by weight of (CF₃)₂CHOCHCl₂.
 - 14. The method of claim 1, wherein the mixture comprises water.
- 15. The method of claim 14, wherein the mixture comprises 1 to 25 percent by weight of water.
 - 16. The method of claim 14, wherein the mixture comprises 1 to 15 percent by weight of water.
- 17. The method of claim 14, wherein the mixture comprises 3 to 10 percent by weight of water.
- 18. The method of claim 1, wherein the mole ratio of the (CF₃)₂CHOCH₂Cl to the hydrogen fluoride is from 1:1 to 1:2.

- 19. The method of claim 1, wherein the mole ratio of the (CF₃)₂CHOCH₂Cl to the amine is from 1:0.3 to 1:2.
- 25 20. The method of claim 3, wherein the mixture is reacted for 4 to 12 hours.
 - 21. The method of claim 3, wherein the mixture is reacted for 4 to 10 hours.
 - 22. The method of claim 3, wherein the mixture is reacted for 4 to 7 hours.
- 23. The method of claim 1, wherein the yield of the reaction is at least 50 percent.

- 24. The method of claim 1, wherein the yield of the reaction is at least 65 percent.
- 25. The method of claim 1, wherein the yield of the reaction is at least 75 percent.
 - 26. The method of claim 1, wherein the conversion of the reaction is at least 50 percent.
- 10 27. The method of claim 1, wherein the conversion of the reaction is at least 60 percent.
 - 28. The method of claim 1, wherein the conversion of the reaction is at least 70 percent.
- 29. The method of claim 1, comprising separating the (CF₃)₂CHOCH₂F from the mixture after the mixture has reacted by washing the mixture with water.
- 20 30. The method of claim 1, comprising separating the (CF₃)₂CHOCH₂F from the mixture by steam distillation.
 - 31. The method of claim 1, comprising separating the (CF₃)₂CHOCH₂F from the mixture by fractional distillation.
 - 32. The method of claim 1, comprising separating the (CF₃)₂CHOCH₂F from the mixture by distillation with an inert high boiling solvent having a boiling point above that of sevoflurane.
- 33. The method of claim 32, wherein the high boiling solvent is an aromatic hydrocarbon.
 - 34. The method of claim 33, wherein the aromatic hydrocarbon is xylene.

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- 35. The method of claim 32, wherein the distillation with the inert high boiling solvent proceeds simultaneously with reacting the mixture.
- 5 36. A method for the preparation of sevoflurane which comprises:
 - (a) providing (CF₃)₂CHOCH₃;

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- (b) contacting the (CF₃)₂CHOCH₃ with chlorine gas;
- (c) exposing the mixture to a sufficient amount of light to initiate and sustain an exothermic reaction;
 - (d) allowing the exothermic reaction to proceed to produce a (CF₃)₂CHOCH₂Cl intermediate; and
 - (e) reacting in a liquid mixture the (CF₃)₂CHOCH₂Cl intermediate with hydrogen fluoride and an amine; to form (CF₃)₂CHOCH₂F.
 - 37. The method of claim 36, wherein the (CF₃)₂CHOCH₂Cl intermediate is reacted with the amine and the hydrogen fluoride by heating.
 - 38. The method of claim 36, wherein the (CF₃)₂CHOCH₂Cl intermediate is reacted with the amine and the hydrogen fluoride by heating at 40°C to 80°C.
- 39. The method of claim 36, wherein the (CF₃)₂CHOCH₂Cl intermediate is reacted with the amine and the hydrogen fluoride by heating at 55°C to 65°C.
 - 40. The method of claim 36, wherein the light is ultraviolet light.
 - 41. The method of claim 36, wherein the amine is a tertiary amine.
- 42. The method of claim 41, wherein the tertiary amine is selected from triethylamine, tripropylamine, triisopropylamine, tributylamine, dimethyl ethyl amine, di-isopropyl ethyl amine, N-methyl pyrrolidine, or mixtures thereof.